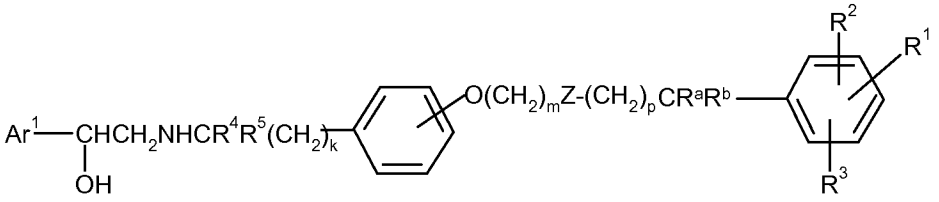
Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

What is claimed is:

1. (Original) A compound of formula (I)



(I)

or a salt, solvate, or physiologically functional derivative thereof, wherein:
k is an integer of from 1 to 3;
m is an integer of from 2 to 4;
p is an integer of from 0 to 3;
Z is O or CH_2-

R^1 is selected from hydrogen, C_{1-6}alkyl, hydroxy, C_{1-6}alkoxy, cyano, nitro, halo, C_{1-6}haloalkyl, XCO_2R^8, $-XC(O)NR^7R^8$, $-XNR^6C(O)R^7$, $-XNR^6C(O)NR^7R^8$, $-XNR^6C(O)NC(O)NR^7R^8$, $-XNR^6SO_2R^7$, $-XSO_2NR^9R^{10}$, XSR^6, $XSOR^6$, XSO_2R^6, $XNR^6SO_2NR^7R^8$, $XNR^6SO_2NR^7COOR^7$,
$-XNR^7R^8$, $-XNR^6C(O)OR^7$,
or R^1 is selected from -X-aryl, -X-hetaryl, or -X-(aryloxy), each optionally substituted by 1 or 2 groups independently selected from hydroxy, C_{1-6}alkoxy, halo, C_{1-6}alkyl,
C_{1-6}haloalkyl, $-NR^6C(O)R^7$, SR^6, SOR^6, $-SO_2R^6$, $-SO_2NR^9R^{10}$, $-CO_2R^8$, $-N\,R^7R^8$, or hetaryl optionally substituted by 1 or 2 groups independently selected from hydroxy, C_{1-6}alkoxy, halo, C_{1-6}alkyl, or C_{1-6}haloalkyl;

X is $-(CH_2)_q$- or C_{2-6} alkenylene;

q is an integer from 0 to 6;

R^6 and R^7 are independently selected from hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aryl, hetaryl, hetaryl(C_{1-6}alkyl)- and aryl(C_{1-6}alkyl)- and R^6 and R^7 are each independently optionally substituted by 1 or 2 groups independently selected from halo, C_{1-6}alkyl,
C_{3-7} cycloalkyl, C_{1-6} alkoxy, C_{1-6}haloalkyl, -NHC(O)(C_{1-6}alkyl), -SO_2(C_{1-6}alkyl), -SO_2(aryl), -CO_2H, and -CO_2(C_{1-4}alkyl), -NH_2, -NH(C_{1-6}alkyl), aryl(C_{1-6}alkyl)-, aryl(C_{2-6}alkenyl)-,
aryl(C_{2-6}alkynyl)-, hetaryl(C_{1-6}alkyl)-, -$NHSO_2$aryl, -NH(hetarylC_{1-6}alkyl), -$NHSO_2$hetaryl,
-$NHSO_2$(C_{1-6}alkyl), -NHC(O)aryl, or -NHC(O)hetaryl:

R^8 is selected from hydrogen, C_{1-6}alkyl and C_{3-7} cycloalkyl;

or R^7 and R^8, together with the nitrogen atom to which they are bonded, form a 5-, 6- or 7- membered nitrogen – containing ring;

R^9 and R^{10} are independently selected from hydrogen, C_{1-6}alkyl, C_{3-7}cycloalkyl, aryl, hetaryl, hetaryl(C_{1-6}alkyl)- and aryl(C_{1-6}alkyl)-, or R^9 and R^{10}, together with the nitrogen to which they are bonded, form a 5-, 6-, or 7- membered nitrogen containing ring;
and R^9 and R^{10} are each optionally substituted by one or two groups independently selected from halo, C_{1-6}alkyl, and C_{3-7}cycloalkyl, C_{1-6}haloalkyl;

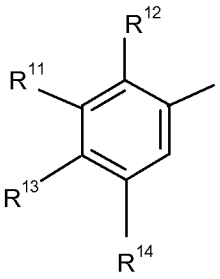
R^2 is selected from hydrogen, hydroxy, C_{1-6}alkyl, C_{1-6}alkoxy, halo, aryl, aryl(C_{1-6}alkyl)-, C_{1-6}haloalkoxy, and C_{1-6}haloalkyl;

R^3 is selected from hydrogen, hydroxy, C_{1-6}alkyl, C_{1-6}alkoxy, halo, aryl, aryl(C_{1-6}alkyl)-, C_{1-6}haloalkoxy, and C_{1-6}haloalkyl;

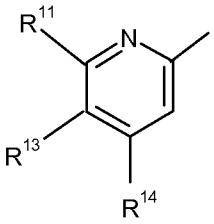
R^a and R^b are independently selected from hydrogen and C_{1-4} alkyl.

R^4 and R^5 are independently selected from hydrogen and C_{1-4} alkyl with the proviso that the total number of carbon atoms in R^4 and R^5 is not more than 4: and

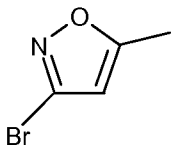
Ar^1 is a group selected from



(a)

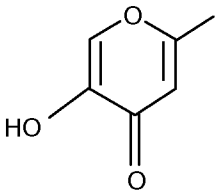


(b)



(c)

and



(d)

wherein R^{11} represents halogen, $-(CH_2)_nOR^{15}$, $-NR^{15}C(O)R^{16}$, $-NR^{15}SO_2R^{16}$, $-SO_2NR^{15}R^{16}$, $-NR^{15}R^{16}$, $-OC(O)R^{17}$ or $OC(O)NR^{15}R^{16}$,
and R^{12} represents hydrogen, halogen or C_{1-4} alkyl;

or R^{11} represents $–NHR^{18}$ and R^{12} and $–NHR^{18}$ together form a 5- or 6-membered heterocyclic ring;

R^{13} represents hydrogen, halogen, $–OR^{15}$ or $–NR^{15}R^{16}$;

R^{14} represents hydrogen, halogen, haloC_{1-4} alkyl, $-OR^{15}$, $–NR^{15}$ R^{16}, $-OC(O)R^{17}$ or $OC(O)NR^{15}R^{16}$;

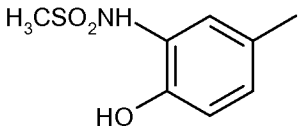
R^{15} and R^{16} each independently represents hydrogen or C_{1-4} alkyl, or in the groups
$-NR^{15}R^{16}$, $-SO_2NR^{15}R^{16}$ and $-OC(O)NR^{15}R^{16}$, R^{15} and R^{16} independently represent hydrogen or C_{1-4} alkyl or together with the nitrogen atom to which they are attached form a 5-, 6- or 7- membered nitrogen-containing ring,

R^{17} represents an aryl group which may be unsubstituted or substituted by one or more substituents selected from halogen, C_{1-4} alkyl,
hydroxy, C_{1-4} alkoxy or halo C_{1-4} alkyl; and

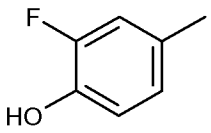
n is zero or an integer from 1 to 4;
provided that in the group (a), when R^{11} represents $-(CH_2)_nOR^{15}$ and n is 1, R^{13} is not OH.

2. (Original) A compound according to claim 1 wherein Ar^1 is selected from group (a) or group (b), as defined in claim 1.

3. (Original) A compound of formula (I) according to claim 2 wherein group (a) is selected from a group of formula (iv) or (xix):

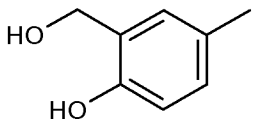


(iv)



(xix)

4. (Original) A compound of formula (I) according to claim 2 wherein group (b) is a group of formula (iii):



(iii)

5. (Currently Amended) A compound of formula (I) according to claim 1 ~~any of claims 1-4~~ wherein R^1 is selected from hydrogen, C_{1-4}alkyl, hydroxy,

cyano, C_{1-6}alkoxy, halo, XCO_2R^8, XNR^6COR^7, $XCONR^7R^8$, $-NR^6C(O)NR^7R^8$, $XSOR^6$, $XNR^6SO_2NR^7R^8$, $XNR^6SO_2NR^7CO_2R^7$ and $-NR^6SO_2R^7$ wherein R^6 and R^7 are as defined above.

6. (Original) A compound of formula (I) according to claim 5 wherein R^1 is selected from $XC(O)NR^7R^8$ or hydrogen.

7. (Currently Amended) A compound of formula (I) according to claim 1 ~~any of claims 1-6~~ wherein R^2 and R^3 each represent hydrogen.

8. (Currently Amended) A compound of formula (I) according to claim 1 ~~any of claims 1-7~~ wherein R^4 and R^5 each represent hydrogen.

9. (Currently Amended) A compound of formula (I) according to claim 1 ~~any of claims 1-8~~ wherein R^a and R^b each represent hydrogen.

10. (Currently Amended) A compound of formula (I) according to claim 1 which is selected from the group consisting of:
3-{[2-(4-{2-[((2*R*)-2-hydroxy-2-{4-hydroxy-3-[(methylsulfonyl)amino]phenyl}ethyl)amino]ethyl}phenoxy)ethoxy]methyl}benzamide;
N-{2-hydroxy-5-[(1*R*)-1-hydroxy-2-({2-[4-(4-phenylbutoxy)phenyl]ethyl}amino)ethyl]phenyl}methanesulfonamide;
N-(5-{(1*R*)-2-[(2-{4-[2-(benzyloxy)ethoxy]phenyl}ethyl)amino]-1-hydroxyethyl}-2-hydroxyphenyl)methanesulfonamide;
3-({2-[4-(2-{[(2*R*)-2-(3-fluoro-4-hydroxyphenyl)-2-hydroxyethyl]amino}ethyl)phenoxy]ethoxy}methyl)benzamide;
4-{(1*R*)-2-[(2-{4-[2-(benzyloxy)ethoxy]phenyl}ethyl)amino]-1-hydroxyethyl}-2-fluorophenol;
2-fluoro-4-[(1*R*)-1-hydroxy-2-({2-[4-(4-phenylbutoxy)phenyl]ethyl}amino)ethyl]phenol;
3-[(2-{4-[2-({2-hydroxy-2-[5-hydroxy-6-(hydroxymethyl)pyridin-2-yl]ethyl}amino)ethyl]phenoxy}ethoxy)methyl]benzamide;

6-{2-[(2-{4-[2-(benzyloxy)ethoxy]phenyl}ethyl)amino]-1-hydroxyethyl}-2-(hydroxymethyl)pyridin-3-ol;
2-(hydroxymethyl)-6-[1-hydroxy-2-({2-[4-(4-phenylbutoxy)phenyl]ethyl}amino)ethyl]pyridin-3-ol;

~~and~~ salts thereof, solvates thereof and physiologically functional derivatives thereof.

11. (Currently Amended) A method for the prophylaxis or treatment of a clinical condition in a mammal~~, such as a human~~, for which a selective β_2-adrenoreceptor agonist is indicated, which comprises administering ~~administration~~ of a therapeutically effective amount of a compound of formula (I), according to claim 1 ~~any of claims 1-10~~, or a pharmaceutically acceptable salt, solvate, or physiologically functional derivative thereof.

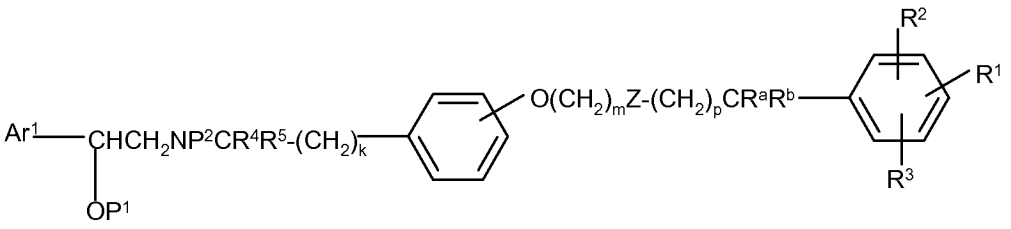
12-13 (Canceled)

14. (Currently Amended) A pharmaceutical formulation comprising a compound of formula (I), according to claim 1 ~~any of claims 1-1~~0, or a pharmaceutically acceptable salt, solvate, or physiologically functional derivative thereof, and a pharmaceutically acceptable carrier or excipient, and optionally one or more other therapeutic ingredients.

15. (Canceled)

16. (Currently Amended) A process for the preparation of a compound of formula (I), according to claim 1 ~~any of claims 1-10~~, or a salt, solvate, or physiologically functional derivative thereof, which comprises:

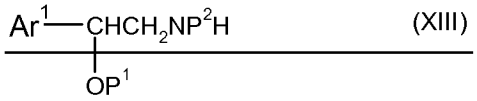
~~(a)~~ ~~deprotection of~~ deprotecting a protected intermediate~~, for example~~ of formula (II):



(II)

or a salt or solvate thereof, wherein Ar^1, R^1, R^2, R^3, R^a, R^b, R^4, R^5, Z, k, m, and p are as defined for the compounds of formula (I), and P^1 and P^2 are each independently either hydrogen or a protecting group provided that at least one of P^1 and P^2 is a protecting group~~; or~~

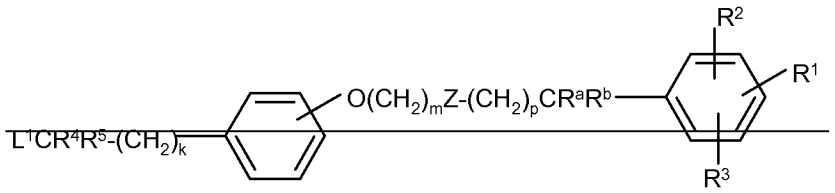
~~(b) alkylation of an amine of formula (XIII)~~



(XIII)

~~wherein Ar^1 is as defined above for compounds of formula (I) and P^1 and P^2 are each independently either hydrogen or a protecting group,~~

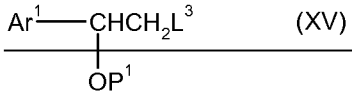
~~with a compound of formula (XIV):~~



(XIV)

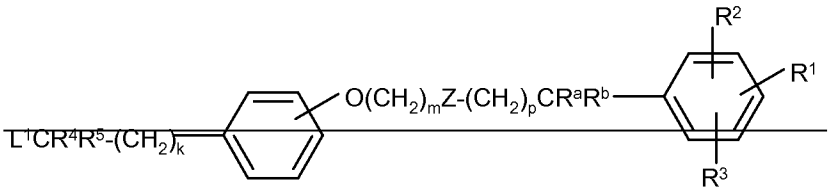
~~wherein R^1, R^2, R^3, R^4, R^5, R^a, R^b, Z, m, and p are as defined for the compound of formula (I) and L^1 is a leaving group;~~

~~(c) reacting a compound of formula (XV):~~



(XV)

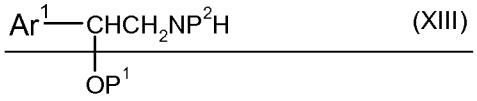
~~wherein P^1 and Ar^1 are as hereinbefore defined and L^3 is a leaving group, with an amine of formula (XVI):~~



(XIV)

~~wherein R^1, R^2, R^3, R^4, R^5, R^a, R^b, Z k, m, p and P^2 are as hereinbefore defined; or~~

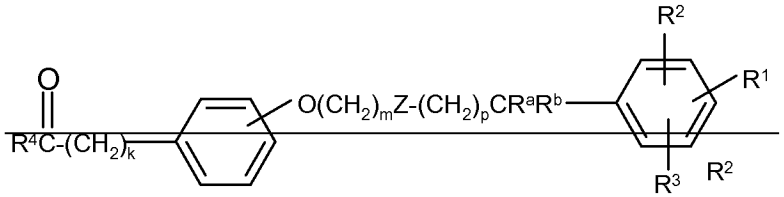
~~d) reacting a compound of formula (XIII):~~



(XIII)

~~as hereinbefore defined,~~

~~with a compound of formula (XVII):~~



(XVII)

~~under conditions suitable to effect reductive amination.~~

wherein said deprotecting step is optionally followed by one or more of the following steps in any order selected from the group consisting of ~~in any order~~:

(i) ~~optional removal of~~ removing any protecting groups;

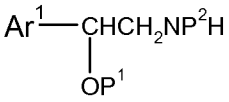
(ii) ~~optional separation of~~ separating an enantiomer from a mixture of enantiomers; and

(iii) ~~optional conversion of~~ converting the product to a corresponding salt, solvate,

or physiologically functional derivative thereof.

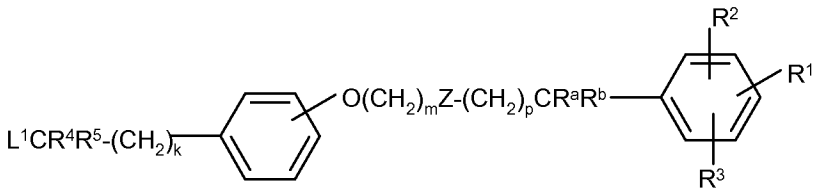
17. (New) A process for the preparation of a compound of formula (I), according to claim 1, or a salt, solvate, or physiologically functional derivative thereof, which comprises:

alkylating an amine of formula (XIII)



(XIII)

wherein Ar^1 is as defined above for compounds of formula (I) and P^1 and P^2 are each independently either hydrogen or a protecting group,
with a compound of formula (XIV):



(XIV)

wherein R^1, R^2, R^3, R^4, R^5, R^a, R^b, Z, m, and p are as defined for the compound of formula (I) and L^1 is a leaving group;

wherein said alkylating step is optionally followed by one or more of the following steps in any order selected from the group consisting of:

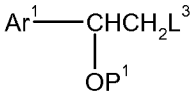
(i) removing any protecting groups;

(ii) separating an enantiomer from a mixture of enantiomers; and

(iii) converting the product to a corresponding salt, solvate,
or physiologically functional derivative thereof.

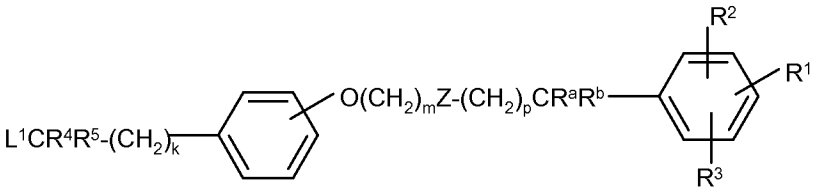
18. (New) A process for the preparation of a compound of formula (I), according to claim 1, or a salt, solvate, or physiologically functional derivative thereof, which comprises:

reacting a compound of formula (XV):



(XV)

wherein P^1 is either hydrogen or a protecting group and Ar^1 are as hereinbefore defined and L^3 is a leaving group, with an amine of formula (XVI):



(XIV)

wherein R^1, R^2, R^3, R^4, R^5, R^a, R^b, Z k, m, p and P^2 are as hereinbefore defined, and L^1 is a leaving group;

wherein said reacting step is optionally followed by one or more of the following steps in any order selected from the group consisting of:

(i) removing any protecting groups;

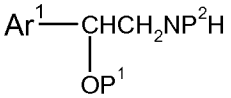
(ii) separating an enantiomer from a mixture of enantiomers; and

(iii) converting the product to a corresponding salt, solvate,

or physiologically functional derivative thereof.

19. (New) A process for the preparation of a compound of formula (I), according to claim 1, or a salt, solvate, or physiologically functional derivative thereof, which comprises:

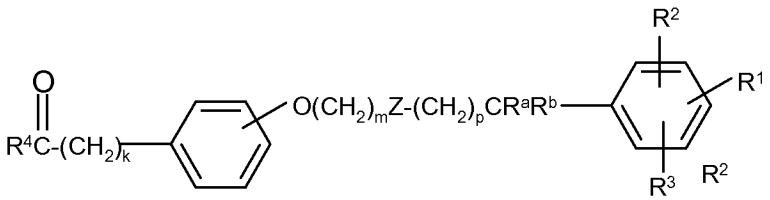
reacting a compound of formula (XIII):



(XIII)

as hereinbefore defined, and wherein P^1 and P^2 are each independently either hydrogen or a protecting group provided that at least one of P^1 and P^2 is a protecting group,

with a compound of formula (XVII):



(XVII)

under conditions suitable to effect reductive amination;

wherein said reacting step is optionally followed by one or more of the following steps in any order selected from the group consisting of:

(i) removing any protecting groups;

(ii) separating an enantiomer from a mixture of enantiomers; and

(iii) converting the product to a corresponding salt, solvate,

or physiologically functional derivative thereof.

20. (New) The method according to Claim 11, wherein said mammal is a human.